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FIRST NAMED INVENTOR CONFIRMATION NO. ATTORNEY DOCKET NO. FILING DATE APPLICATION NO. 8132 600.311USD1 Patrick M. Schlievert 07/22/2003 10/625,221 EXAMINER 05/17/2004 7590 GRASER, JENNIFER E Attention of Mark T. Skoog MERCHANT & GOULD P.C. PAPER NUMBER ART UNIT P.O. Box 2903 1645 Minneapolis, MN 55402-0903

DATE MAILED: 05/17/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)			
·	10/625,221	SCHLIEVERT ET AL.			
Office Action Summary	Examiner	Art Unit			
<i></i>	Jennifer E. Graser	1645			
The MAILING DATE of this communication app	I .				
Period for Reply					
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).					
Status					
1) Responsive to communication(s) filed on	<u>_</u> .				
, :	action is non-final.				
3) Since this application is in condition for allowa	nce except for formal matters, pro	secution as to the merits is			
closed in accordance with the practice under I					
Disposition of Claims					
4)⊠ Claim(s) <u>40-60</u> is/are pending in the applicatio	n.	•			
4a) Of the above claim(s) is/are withdra					
5) Claim(s) is/are allowed.		**			
6)⊠ Claim(s) <u>40-60</u> is/are rejected.					
7) ☐ Claim(s) is/are objected to.					
8) Claim(s) are subject to restriction and/o	or election requirement.				
Application Papers					
9)☐ The specification is objected to by the Examine	er.				
10)⊠ The drawing(s) filed on <u>7/22/03</u> is/are: a)⊠ ad	ccepted or b) \square objected to by the	e Examiner.			
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).					
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).					
11)☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.					
Priority under 35 U.S.C. § 119					
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received.					
Attachment(s) 1) Notice of References Cited (PTO-892)	4)	/ (PTO-413) late.			
2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08		Patent Application (PTO-152)			
Paper No(s)/Mail Date 10/24/03.	6) Other:				

Art Unit: 1645

DETAILED ACTION

Priority

1. This application filed under former 37 CFR 1.60 lacks the necessary reference to the prior application. A statement reading "This is a Divisional of Application No. 08,973,391, now U.S. Patent No. X, filed 3/12/98, which is a CIP of....." should be entered following the title of the invention or as the first sentence of the specification. Also, the current status of all nonprovisional parent applications referenced should be included.

Specification

2. The disclosure is objected to because of the following informalities:

In the "Brief Description of the Drawings", "Figure 4", "Figure 5", and "Figure 6" must be changed to "Figure 4A and 4B", "Figure 5A and 5B", and Figure "6A and 6B" so that they properly correspond to the Drawings.

Appropriate correction is required.

Claim Rejections - 35 USC § 112

- 3. The following is a quotation of the second paragraph of 35 U.S.C. 112:
 The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.
- 4. Claims 40-60 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claims 40-44, 50, 52, and 54 are vague and indefinite because the claim fails to teach what the substitution of the amino acid should be substituted with, thereby leaving

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it unclear as the structure of the claimed nucleic acid. Instead, the claims broadly allow for the change to the nucleic acid to encode any amino acid whether it is positively(negatively) charged amino acid, an uncharged amino acid or a hydrophobic(hydrophilic) amino acid. While the specification can be used to provide definitive support, the claims are not read in a vacuum. Rather, the claim must be definite and complete in and of itself. Limitations from the specification will not be read into the claims. The claims as they stand are incomplete and fail to provide adequate structural properties to allow for one to identify what is being claimed. The specific substitution, i.e., asparatine-20 to aspartic acid.

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Claims 56, 59 and 60 are vague and confusing due to the phrases "a polynucleotide having the portion of sequence SEQ ID NO: 12 that encodes a polypeptide having the sequence of SEQ ID NO:14" and "a polynucleotide having 99% sequence identity with the portion of sequence SEQ ID NO: 12 that encodes a polypeptide having the sequence of SEQ ID NO:14". The claims should be amended so that the phrase "portion of sequence SEQ ID NO: 12 that encodes a polypeptide having the sequence of SEQ ID NO:14" positively recites the portion by sequence position number, i.e., a polynucleotide consisting of nucleotides A-D of SEQ ID NO:12 (wherein A-D are the specific nucleotides which encode SEQ ID NO:14). This wording would clarify the claim.

Claim Rejections - 35 USC § 112

5. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the

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art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

6. Claims 40-44, 50, 52, and 54 are rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the enablement requirement. The claim(s) contains subject matter which was not described in the specification in such a way as to enable one skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the invention.

These claims are not enabled because they do not specify the specific amino acid to which the designated SPE-A amino acid is to be changed to. Instead, the claims broadly allow for the isolated nucleic acid molecules to encode any whether it is a positively(negatively) charged amino acid, an uncharged amino acid or a hydrophobic(hydrophilic) amino acid. The specification teaches that the nucleic acid molecules encode polypeptides to be used in compositions, vaccines and methods of providing protection against wild type SPE-A toxin comprising this breadth given the unpredictability set forth in the Examples provided in the instant specification and the examples taught in the prior art.

The specification and prior art both teach that the results from the substitution of amino acids in the SPE-A mutant is very unpredictable. To change a charged amino acid to an uncharged amino acid changes the chemical nature of the compound which has been shown to negatively impact its function. The specification has shown that substitutions of amino acids, even with the same charge, can cause the resultant SPE-A to lose immunogenicity and it's ability to provide immune protection. Due to the highly unpredictable nature of determining acceptable substitutions, it has been necessary to

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provide specific results and direction as to what the changes may be. Applicants provided these results in the Declaration of Dr. Patrick M. Shlievert submitted in parent case 08/973,391. However, claims which allow for open-ended changes, i.e., that do not recite what the specific substitution can be, are not enabled. There is no showing in the instant specification of the consequences of these broad mutations or their ability to produce a polypeptide which would provide an immune effect in a host. Both the prior art and specification have shown that even when an amino acid of SPE-A toxin is replaced with an amino acid of the same charged, i.e., conservative substitution, it does not guarantee that the SPE-A toxin will retain it's function. The prior art and specification have shown in several instances that such substitutions have resulted in a toxin with reduced immune function. There is also no evidence provided in the specification which indicates whether or not the specific mutants actually have the various claimed properties, i.e., nonlethality, decrease in mitogenicity, not enhancing endotoxin shock, etc.. The number of mutant SPE-A toxins to be encoded by the claimed nucleic acids encompassed by the instant claims is vast and would not allow one of skill in the art a reasonable expectation of success that any mutant encompassed by the instant claims would have the claimed properties. Specific quidance is needed for the skilled artisan to make a reasoned decision of which mutants would likely have the most success in producing polypeptides which would be effective in vaccine compositions. It is unpredictable as to which amino acids could be removed and which could be added. While it is known that many amino acid substitutions are possible in any given protein, the position within the protein's sequence where amino

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acid substitutions can be made with a reasonable expectation of success are limited. Other positions are critical to the protein's structure/function relationship, e.g., such as various positions or regions directly involved in binding, catalysis in providing the correct three-dimensional spatial orientation of binding and catalytic sites. These regions can tolerate only very little or no substitutions. To start with the DNA sequence first, this requires even more work on the part of the skilled artisan. Applicants have not provided sufficient guidance to enable one of ordinary skill in the art how to determine, without undue experimentation, the effects of different amino acid substitutions and the nature and extent of the changes that can be made.

Given the lack of guidance contained in the specification and the unpredictability for determining acceptable nucleotide substitutions and their effect on the immunogenicity of the SPE-A toxin they encode, one of skill in the art could not make or use the broadly claimed invention without undue experimentation.

Allowable Subject Matter

- 7. Claims 45-49, 51, 53, 55, and 56-60 would be allowable if rewritten to overcome the rejection(s) under 35 U.S.C. 112, second and first paragraph, set forth in this Office action and to include all of the limitations of the base claim and any intervening claims.
- 8. Correspondence regarding this application should be directed to Group Art Unit 1645. Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Group 1600 via the PTO Fax Center located in Remsen. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15,1989). The Group 1645 Fax number is (703) 872-9306 which is able to receive transmissions 24 hours/day, 7 days/week.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jennifer E. Graser whose telephone number is (571)

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272-0858. The examiner can normally be reached on Monday-Friday from 7:00 AM-4:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Lynette Smith, can be reached on (571) 272-0864.

Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist whose telephone number is (571) 272-0500.

ennifer Graser

Primary Examiner

Art Unit 1645

PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:

SCHLIEVERT ET AL.

Examiner:

UNKNOWN

Serial No.:

10/625,221

Group Art Unit:

1645

Filed:

JULY 22, 2003

Docket No.:

600.311USD1

Confirmation No.:

UNKNOWN

Customer No.:

23552

Title:

MUTANTS OF STREPTOCOCCAL TOXIN A AND METHOD OF USE

CERTIFICATE UNDER 37 CFR 1.8:

Thereby certify that this correspondence is being deposited with the United States Postal Service as first class mail, with sufficient postage, in an envelope addressed to: Commissioner for Patents, P.O. Box 450, Alexandria, VA 22313-1450 on October 22, 2003.

Name: Shervi A. Boerboom

INFORMATION DISCLOSURE STATEMENT (37 C.F.R. § 1.97(b))

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

With regard to the above-identified application, the items of information listed on the enclosed Form 1449 are brought to the attention of the Examiner.

This statement should be considered because it is submitted within three months of the filing date of the above-identified application, which is not an application under 37 C.F.R. § 1.53(d). Accordingly, no fee is due for consideration of the items listed on the enclosed Form 1449.

In accordance with 37 C.F.R. §1.98(d), a copy of each document or other information listed on the enclosed Form 1449 is not provided because it was previously cited by or submitted to the U.S. Patent and Trademark Office in parent application, U.S. Serial No. 08/973,391 filed on March 12, 1998.

No representation is made that a reference is "prior art" within the meaning of 35 U.S.C. §§ 102 and 103 and Applicants reserve the right, pursuant to 37 C.F.R. § 1.131 or otherwise, to establish that the reference(s) are not "prior art." Moreover, Applicants do not represent that a

reference has been thoroughly reviewed or that any relevance of any portion of a reference is intended.

Consideration of the items listed is respectfully requested. Pursuant to the provisions of M.P.E.P. 609, it is requested that the Examiner return a copy of the attached Form 1449, marked as being considered and initialed by the Examiner, to the undersigned with the next official communication.

Please charge any additional fees or credit any overpayment to Deposit Account No. 13-2725.

Respectfully submitted,

MERCHANT & GOULD P.C. P.O. Box 2903 Minneapolis, MN 55402-0903 (612) 332-5300

DATE: Oct 22, 2003

Mark T. Skoog Reg. No. 40,178

PATENT TRADEMARK OFFICE

EOIDHIA.	INFORMATION DISCLOSURE STATEMENT	Docket Number: 600.311USD1	Application Number: 10/625,221	
OCI 2 (2003 2)	IN AN APPLICATION	IN AN APPLICATION Applicant: SCHLIEVERT ET AL.		
	(Use several sheets if necessary)	Filing Date: July 22, 2003	Group Art Unit: 1645	
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7	7	5,298,396		03/29/1994	Kotzin et al.				
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	A	WO 93/1463	4	08/05/1993	PCT				
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	\sim		ОТІ	HER DOCUMEN	TS (Including Author, Title, Da	ate, Pertinent Pages,	Etc.)		
			Acharya, K. et al., "Structural Basis of Superantigen Action Inferred from Crystal Structure of Toxic-Shock Syndrome Toxin-1", Nature 367:94-97 (1994).					`oxin-l",	
T			Aiyar, A. et al., "Modification of the Megaprimer Method of PCR Mutagenesis: Improved Amplification of the Final Product", BioTechniques Vol. 14, No. 3 (1993) pages 366-369.						
			Altschyl, S. e	Altschyl, S. et al., "Optimal Sequence Alignment Using Affine Gap Costs", Bulletin of Math. Biol. 48:603-616 (1986).					
			Anthony-Cahil, S. et al., "Site-specific mutagenesis with unnatural amino acids", Trends in Biochem. Sci. 14:400-403 (1989).						
			Barsumian et al., "Nonspecific and Specific Immunological Mitogenicity by Group A Streptococcal Pyrogenic Exotoxins", Infection and Immunity 22:681-688 (1978).						
			Belani, K. et al., "Association of exotoxin-producing Group A streptococci and severe disease in children, Pediatr. Infect. Dis. J. 10:351-354 (1991).						
			Betley et al., "Staphylcoccal Enterotoxins, Toxic Shock Syndrome Toxin and Streptococcal Pyrogenic Exotoxins: A Comparative Study of their Molecuar Biology", Chem. Immun. 55:1-35 (1992).						
			Birkhaug et al., "Studies in Scarlet Fever II: Studies on the Use of Convalescent Scarlet Fever Serum in Dochez Scarletino Antistreptococcic serum for the treatment of scarlet fever", Bull. John Hopkins Hosp. 36:134-171 (1925).						
			Black, C.M. et al., "Detection of Streptococcal Pyrogenic Exotoxin Genes by a Nested Polymerase Chain Reaction", Molecular and Cellular Probes, Vol. 7, pp. 255-259 (1993).						
-			Bohach et al., "Staphylcoccal and Streptococcal Pyrogenic Toxins Involved in Toxic Shock Syndrome and Related Illnesses", Crit. Rev. Microbiol. 17:251-272 (1989).						
			Bowie, J. et al., "Deciphering the Message in Protein Sequences: Tolerance to Amino Acid Substitutions", Science 247:1306-1310 (March 16, 1990).						
					es in Both Class II Major Histor drome Toxin I", J. Exper. Med			ntribute to the B	inding of the
			Dohlsten et al., "Superantigen Induced Cytokines Supress Growth of Human Colon Carcinoma Cells", Int. J. Cancer 54:482-488 (1993).				182-488		
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EXAMINER

DATE CONSIDERED

EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form for next communication to the Applicant.

Application Number:

FORM HE	ANDORSE STONE DECOLOGING OF A TOMPSON	Docket Number:	Application Number:			
IN AN APPLICATION		600.311USD1	10/625,221			
		Applicant: SCHLIEVERT ET AL.				
100.	(Use several sheets if necessary)	Filing Date: July 22, 2003	Group Art Unit: 1645			
TO TRACTURE						
	Fast, D. et al., "Toxic Shock Syndrome-Associated St. Necrosis Factor Production", Infection and Immunity		ic Toxins are Potent Inducers of Tumor			
	Goshorn, S. et al., "Cloning and characterization of the Gen. Genet. 212:66-70 (1988).	the gene, speC, for pyrogenic exotoxin type C from Streptococcus pyogenes", Mol.				
	Goshom, S. et al., "Nucelotide Sequence of Streptoco	Goshom, S. et al., "Nucelotide Sequence of Streptococcal Pyrogenic Exotoxin Type C", Infection and Immunity 56:2518-2520 (1988)				
	Griggs, N. et al., "Mapping of Multiple Binding Dome 148:2516-2521 (April 15, 1992).	Griggs, N. et al., "Mapping of Multiple Binding Domains of the Superantigen Staphylococcal Enterotoxin A for HLA", J. Immunolog 148:2516-2521 (April 15, 1992).				
	Hartwig, U. et al., 1993. "Mutations affecting MHC of International Immunology 5(8):869-875.	Hartwig, U. et al., 1993. "Mutations affecting MHC class II binding of the superantigen streptococcal erythrogenic toxin A." International Immunology 5(8):869-875.				
	Hattori, M. et al., "Structure of the rat α2-macroglobulin-coding gene", Gene 77:333-340 (1989).					
		Hauser, A. et al., "Molecular Analysis of Pyrogenic Exotoxins from Streptococcus pyogenes Isolates Associated with Toxic Shock-Like Syndrome", J. Clin. Microbiol. 29:1562-1567 (August 1991).				
	Hedlund et al., "Superantigen-Based Tumor Therapy in Vivo Activation of Cytotoxic T Cells", Cancer Immun. Immunother. 36:89-(1993).					
	Hsiao, Ku-chuan et al., "A Fast and simple procedure 19:2787 (1991).	Hsiao, Ku-chuan et al., "A Fast and simple procedure for sequencing double stranded DNA with Sequence", Nucleic Acids Resea 19:2787 (1991).				
	Ihle et al., "Antibody Targeted Super Antigens Induce Lines", Cancer Res. 55:623-628 (1995).	Ihle et al., "Antibody Targeted Super Antigens Induce Lysis of Major Histocompatibility Complex Class II Negative T Cell Leuken Lines", Cancer Res. 55:623-628 (1995).				
	lwasaki et al., "Cloning, Characterization and Overext Factor", FEBS Lett. 331:187-192 (1993).	ion and Overexpression of Streptococcus Pyogenes Gene Encoding a New Type of Mitogenic 3).				
	Jardetzky, T. et al., "Three-dimensional structure of a Nature 368:711-718 (April 21, 1994).	sional structure of a human class II histocompatibility molecule complexed with superantigen", 94).				
		leoccal Enerotoxin B Sequences Important for Induction of Lymphocyte Proliferation Using Toxin", Infection and Immunity 62:3408-3415 (1994).				
	Johnson, L. et al., "Group A streptococcal phage T12 (194:52-56 (1994).	o A streptococcal phage T12 carries the structural gene for pyrogenic exotoxin type A", Mol. Gen. Genet.				
	Kappler, J. et al., "Mutations Defining Functional Reg 396 (February 1992).					
	Lee, P. et al, "Effects of Staphylococcal Toxic Shock Syndrome Toxin 1 on Aortic Endothelial Cells", J. Infect. Dis. 164:711-9 (1991).					
	Lee, P. et al., "Fluid Replacement Protection of Rabbit and Immunity 59(3):879-884 (Mar 1991).	ts Challenged Subcutanteously with To	xic Shock Syndrome Toxins", Infection			
	Marrack, P. et al., "The Staphylcoccal Enterotoxins and Their Relatives", Science 248:705-711 (May 1990).					
	Martin, D., et al., "Molecular Epidemiology of Group	A Streptococcus M Type 1 Infections, I	l. Infect. Dis. 167;1112-7 (1993).			
	Mollick, J. et al., "Localization of a Site on Bacterial S Med. 177:283-293 (February 1993).	Superantigens That Determines T Cell	Receptor β Chain Specificity*, J. Exp.			
	Mollick, J. et al., "Novel Superantigen Isolated from P. Staphylcoccal Enterotoxins B and C", J. Clin. Invest. S		genes with Aminoterminal Homology to			
	Murray, D. et al., "Immunobiologic and Biochemical (1984) 152(1):87-95.	Properties of Mutants of Toxic Shock S	Syndrome Toxin-1", J. Immunol (US)			
		- 1	3/12/			

DATE CONSIDERED

EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form for next communication to the Applicant.

EXAMINER

INFORMATION DISCLOSURE STATEMENT

Application Number:

Docket Number:

TIPE !	INFORMAT	FION DISCLOSURE STATEMENT	600.311USD1	10/625,221	
	IN AN APPLICATION (Use several sheets if necessary)		Applicant: SCHLIEVERT ET AL.		
oci s t ann			Filing Date: July 22, 2003	Group Art Unit: 1645	
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0		Musser et al., "Streptococcus Pyogenes Causing Pyrogenic Exotoxin Expression", Proc. Nat'l. Ac	Toxic Shock-like Syndrome and Other Ir ad. Sci. (USA) 88:2668-2672 (1991).	vasive Diseases: Colonal Diversity and	
		Musser, J. et al., "Infect Immun", Mar. 1995, 63	(3) P994-1003		
		Nelson, K. et al., "Characterization and Clonal Distribution of Four Alleles of the speA Gene Encoding pyrogenic Exotoxin A (Scarle Fever Toxin) in Streptococcus pyogenes", . Exp. Med., 174:1271-1274 (Nov. 1991) Norrby-Teglund, A. et al., "Relation between Low Capacity of Human Sera to Inhibit Streptococcal Mitogens and Serious Manifestation of Disease", J. Infect. Dis. 170:585-91 (1994). Perrin, S. et al., "Site-specific mutagenesis using asymmetric polymerase chain reaction and a single mutant primer", Nucleic Acids Research 18:7433-7438 (1990).			
		Prasad, G. et al., "Structure of Toxic Shock Syndrome Toxine 1", Biochemistry Vol. 32, No. 50 (December 21, 1993) 50:13761-13766.			
		Rennell, D. et al, "Systematic Mutation of Bacteriophage T4 Lysozyme", J. Mol. Biol. 222:67-87 (1991).			
		Revie, D., et al., "Kinetic analysis for optimization of DNA ligation reactions", Nucleic Acids Research 16:10301-10321 (1988). Roggiani, A. et al., "Localization of biological activities of Streptococcal Pyrogenic Exotoxin", poster presentation at the ASM 94th General Meeting, Las Vegas, Nevada (1994).			
		Schlievert et al., "Group B Streptococcal Toxic S Toxin", Clin. Infect. Dis. 17:26-31 (1993).	lievert et al., "Group B Streptococcal Toxic Shock-Like Syndrome: Report of a Case and Purification of Associated Pyrogenic in", Clin. Infect. Dis. 17:26-31 (1993).		
		Schlievert, "Role of Superantigens in Human Disease", J. Infect. Dis. 167:997-1002 (1993). Schlievert, P. et al., "Infect Immun", June 1989, 57 (6) P1865-7			
		Scott et al., "Characterization of Staphylcoccus a Infection Chambers in Rabbits, Infection and Im	nylcoccus aureus Isolates from Patients with Toxic Shock Syndrome, Using Polyethylene ion and Immunity 39:383-387 (January 1983).		
		Swaminathan, "Crystal Structure of Staphococcal Enterotoxin B as Superantigen", Nature 359:801-806 (1992).			
		Tomai, M. et al., "Distinct T-Cell Receptor Vβ G Exotoxins and pep M5 Protein", Infection and In		imulated with the Streptococcal Pyrogenic	
		Wallace, C., "Understanding cytochrome c function (1993).	ion: engineering protein structure by sen	isynthesis, FASEB Journal 7:505-515	
		Weeks et al., "Nucleotide Sequence of the Type Bacteriophage T12", Infection and Immunology,		Toxin) Gene from Streptococcus pyogenes	

23552 PATENT TRADEMARK OFFICE

EXAMINER

DATE CONSIDERED

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